Appl. No. 09/834,760 Amdt. dated November 14, 2005 Reply to Office Action of October 12, 2004

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

1-46 (canceled)

1 47 (currently amended) A method of inhibiting the generation of active thrombin 2 on the surface of a cell within an atherosclerotic plaque within a mammal, the method 3 comprising increasing the expression or activity of an ER resident calcium-binding protein in 4 said cell by introducing a polynucleotide operably linked to a promoter into said cell, wherein 5 said polynucleotide encodes said ER resident calcium-binding protein, and wherein said ER 6 resident calcium-binding protein is a member selected from the group consisting of GRP78/BiP, - 7 GRP94, GRP72, Calreticulin, Calnexin, Reticulocalbin, Protein disulfide isomerase, cis/trans-8 Prolyl isomerase, and HSP47. 1 48 (previously presented) The method of claim 47, wherein said cell is an 2 endothelial cell. 1 49 (previously presented): The method of claim 47, wherein said cell is a smooth 2 muscle cell. 1 50 (previously presented): The method of claim 47, wherein said cell is a 2 macrophage. 1 51 (previously presented): The method of claim 47, wherein said cell is a 2 monocyte. 1 52 (previously presented): The method of claim 47, wherein said ER resident 2 calcium-binding protein is GRP78/BiP.

Appl. No. 09/834,760 Amdt. dated August 27, 2003 Reply to Office Action of October 12, 2004

1	53 (previously presented): The method of claim 47, wherein said ER resident
2	calcium-binding protein is selected from the group consisting of GRP94, GRP72, Calreticulin,
3	Calnexin, Reticulocalbin, Protein disulfide isomerase, cis/trans-Prolyl isomerase, and HSP47.
l	54 (previously presented): The method of claim 47, wherein the increase in the
2	expression or activity of said ER resident calcium-binding protein within said cell results in a
3	decrease in the level of tissue factor procoagulant activity on the surface of said cell.
	55 (canceled)
l	56 (previously presented): The method of claim 47, wherein said polynucleotide
2	is introduced into said cell using a viral vector.
]	57 (previously presented): The method of claim 56, wherein said viral vector is
2	an adenoviral vector.
l	58 (previously presented): The method of claim 47, wherein said polynucleotide
2	is introduced into said cell using a nonviral vector.
	59 (previously presented): The method of claim 58, wherein said nonviral vector
2	is introduced into said cell as naked DNA or using liposome-mediated transfection.
	60-61 (canceled)
l	62 (currently amended): A method of inhibiting the generation of active
2	thrombin on the surface of a cell within a mammal, the method comprising increasing the
3	expression or activity of an ER resident calcium-binding protein in said cell by administering a
ļ	pro_inflammatory cytokine to said cell, wherein said pro-inflammatory cytokine is a member
;	selected from the group consisting of interleukin-3 and CSF-1.
	63-66 (canceled)

Appl. No. 09/834,760 Amdt. dated November 14, 2005 Reply to Office Action of October 12, 2004 **PATENT**

- 1 67 (previously presented): The method of claim 62, wherein said
- 2 proinflammatory cytokine is interleukin-3.